



Definition of “Small Gram Quantity Contents” for Type B Radioactive Material Transportation Packages: Activity-Based Content Limitations, Rev. 1

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Background and Introduction

Since the 1960's, the Department of Transportation Specification (DOT Spec) 6M packages have been used extensively for transportation of Type B quantities of radioactive materials between Department of Energy (DOE) facilities, laboratories, and productions sites. However, due to the advancement of packaging technology, the aging of the 6M packages, and variability in the quality of the packages, the DOT implemented a phased elimination of the 6M specification packages (and other DOT Spec packages) in favor of packages certified to meet federal performance requirements.^[1] DOT issued the final rule in the Federal Register on October 1, 2004 requiring that use of the DOT Specification 6M be discontinued as of October 1, 2008. A main driver for the change was due to the fact that the 6M specification packagings were not supported by a Safety Analysis Report for Packaging (SARP) that was compliant with Title 10 of the Code of Federal Regulations part 71 (10 CFR 71).^[2,3]

Therefore, materials that would have historically been shipped in 6M packages are being identified as contents in Type B (and sometimes Type A fissile) package applications and addendums that are to be certified under the requirements of 10 CFR 71. The requirements in 10CFR71 include that the Safety Analysis Report for Packaging (SARP) must identify the maximum radioactivity of radioactive constituents and maximum quantities of fissile constituents (10CFR71.33(b)(1) and 10CFR71.33(b)(2)), and that the application (i.e., SARP submittal or SARP addendum) demonstrate that the external dose rate (due to the *maximum radioactivity of radioactive constituents and maximum quantities of fissile constituents*) on the surface of the packaging (i.e., package and contents) not exceed 200 mrem/hr (10CFR71.35(a), 10CFR71.47(a)).

It has been proposed that a ‘Small Gram Quantity’ of radioactive material be defined, such that, when loaded in a transportation package, the dose rates at external points of an unshielded packaging not exceed the regulatory limits prescribed by 10 CFR 71 for non-exclusive shipments. The mass of each radioisotope presented in this paper is limited by the radiation dose rate on the external surface of the package, which per the regulatory limit should not exceed 200 mrem/hr. The results presented are a compendium of allowable masses (and activities) of a variety of different isotopes (with varying impurity levels of beryllium in some of the actinide isotopes) that, when loaded in an unshielded packaging, do not result in an external dose rate on the surface of the package that exceeds 190 mrem/hr (190mrem/hr was chosen to provide 5% conservatism relative to the regulatory limit). These mass limits *define* the term “Small Gram Quantity” (SGQ) contents in the context of radioactive material transportation packages. The term SGQ is isotope-specific and pertains to contents in radioactive material transportation packages that do not require shielding and still satisfy the external dose rate requirements. Since these calculated mass limits are for contents without shielding, they are conservative for

packaging materials that provide some limited shielding or if the contents are placed into a shielded package.

The isotopes presented in this paper were chosen as the isotopes that Department of Energy (DOE) sites most likely need to ship. Other more rarely shipped isotopes, along with industrial and medical isotopes, are planned to be included in subsequent extensions of this work.

Isotopes Considered

Table 1 provides a list of the isotopes considered in Phase 1. This list of isotopes was developed with input from many groups in the DOE complex and is intended to represent a comprehensive list of the isotopes that would generally need to be shipped for purposes related to DOE activities.

Table 1. Isotopes Considered in Analysis

Am-241	Pu-242	Pu-241 w/Be	Hf-181	Ru-106
Am-243	Cf-252	Pu-242 w/Be	Ho-166	Sc-46
Cm-244	Am-241 w/ Be	Ac-227	Ir-192	Se-75
Cm-248	Am-243 w/Be	Cd-109	Mn-54	Sm-145
Np-237	Cm-244 w/Be	Co-60	Pb-210	Sr-90
Pu-238	Np-237 w/Be	Cs-137	Pd-103	Tm-170
Pu-239	Pu-238 w/Be	Eu-152	Pm-147	Yb-169
Pu-240	Pu-239 w/Be	Fe-59	Po-210	Zn-65
Pu-241	Pu-240 w/Be	Gd-153	Ra-226	Zr-95

Approach

For each of the isotopes considered in this analysis, the source term was determined. Table 2 presents the source terms for the neutron-emitting actinides with various weight percentages of beryllium. Table 3 presents the source terms of the gamma-emitting isotopes. All sources are based on 1g of the radioisotope.

The gamma and neutron source spectra from the neutron emitters were obtained using the computer code RASTA.^[4] The oxide chemical form was assumed for determining the source terms for the neutron-emitting isotopes (Table 2) because this is the form of these isotopes most commonly shipped, and also to account for the (α ,n) reaction that occurs with the natural occurring oxygen-17 and oxygen-18 isotopes. For the isotopes with beryllium impurity, the neutron source from the (α , n) reaction was obtained for weight percentages of beryllium from 0–90%. The source terms for the pure gamma emitters were obtained using the computer code ORIGEN-ARP.^[5]

Two different shielding models were used for the calculations: (1) the voided sphere model, and (2) the 9977-96 radioactive material transportation packaging. The source terms, along with the shielding models, were used to calculate the dose rates with MCNP.^[6] The “voided sphere” model is used to determine limiting isotopic masses with no shielding present and only an air gap between the point source and the point of dose rate determination. A representation of the voided sphere model is given in Figure 1. The distance between the point source and the point of dose rate determination was chosen as 15.6 cm. This distance of 15.6 cm corresponds to the

distance from the inner wall of the containment vessel to the outer drum wall in the Model 9977-96 radioactive material transportation packaging. In this way, the limiting mass results from the voided sphere model can be compared to the limiting masses resulting from the 9977 transportation packaging model. A diagram of the Model 9977-96 packaging is shown in Figure 2. A graphical representation of the shielding model used in the MCNP calculations for the 9977-96 package is depicted in Figure 3. The DOE Certificate of Compliance for the Model 9977-96 provides further details concerning the packaging.^[7,8]

Using the source terms in Tables 2 and 3 and the shielding models described, the mass of each isotope in Table 1 that results in a dose rate at the point of dose rate determination of 190 mrem/hr was calculated using the Monte Carlo radiation transport computer code MCNP for both the voided sphere model and for the Model 9977-96. The sources were treated as point sources in the simulation though it is shown as a sphere of finite volume in Figure 2 for the purposes of clarity. For the Model 9977-96, the limiting dose rate occurred on the external side surface of the package, which corresponds to the distance 15.6 cm. These mass limits meet all requirements for non-exclusive use shipments per 10 CFR Part 71.

Table 2. Source Terms for Neutron-Emitting Isotopes

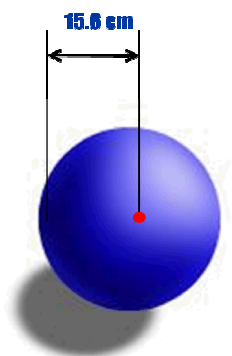
Source Strength [Particles per Second] for One Gram of Actinide Isotope											
Isotope	Gamma	Neutron									
		Weight Percent Beryllium									
		0	0.1	0.5	1	5	10	30	50	70	90
Am-241	1.47E+11	2.76E+03	2.64E+04	1.20E+05	2.35E+05	1.06E+06	2.09E+06	5.05E+06	7.05E+06	8.49E+06	9.58E+06
Am-243	1.96E+10	1.42E+02	NC	NC	NC	NC	1.07E+05	2.57E+05	3.57E+05	4.30E+05	4.84E+05
Cf-252	9.61E+12	2.31E+12	NA	NA	NA	NA	NA	NA	NA	NA	NA
Cm-244	3.18E+11	1.09E+07	1.16E+07	1.43E+07	1.76E+07	4.17E+07	7.14E+07	1.57E+08	2.14E+08	2.56E+08	2.87E+08
Cm-248	2.20E+08	4.03E+07	NA	NA	NA	NA	NA	NA	NA	NA	NA
Np-237	7.09E+07	3.52E-01	NC	NC	NC	NC	2.61E+02	6.31E+02	8.80E+02	1.06E+03	1.20E+03
Pu-238	7.15E+10	1.65E+04	1.35E+05	6.01E+05	1.17E+06	5.06E+06	1.04E+07	2.52E+07	3.53E+07	4.25E+07	4.80E+07
Pu-239	9.80E+07	3.96E+01	3.83E+02	1.74E+03	3.40E+03	1.54E+04	3.03E+04	7.31E+04	1.02E+05	1.23E+05	1.38E+05
Pu-240	9.04E+08	1.17E+03	2.44E+03	7.46E+03	1.36E+04	5.80E+04	1.13E+05	2.71E+05	3.77E+05	4.54E+05	5.11E+05
Pu-241	1.31E+11	2.46E+03	NC	NC	NC	NC	1.86E+06	4.50E+06	6.28E+06	7.56E+06	8.53E+06
Pu-242	1.31E+07	1.72E+03	1.74E+03	1.81E+03	1.90E+03	2.54E+03	3.33E+03	5.58E+03	7.09E+03	8.16E+03	8.97E+03

NC=Not Calculated

NA=Not Applicable

Table 3. Source Terms for Gamma-Emitting Isotopes

Source Strength (Photons per Second) for One Gram of Isotopic Source			
Isotope	[Gammas per Second]	Isotope	[Gammas per Second]
Ac-227	8.41E+12	Pm-147	7.52E+11
Cd-109	8.95E+13	Po-210	1.75E+09
Co-60	8.56E+13	Ra-226	1.25E+11
Cs-137	3.25E+12	Ru-106	1.93E+14
Eu-152	1.48E+13	Sc-46	2.42E+15
Fe-59	1.93E+15	Se-75	1.09E+15
Gd-153	2.23E+14	Sm-145	1.31E+14
Hf-181	1.15E+15	Sr-90	1.78E+12
Ho-166	1.45E+16	Tm-170	4.45E+13
Ir-192	8.12E+14	Yb-169	3.10E+15
Mn-54	2.66E+14	Zn-65	1.48E+14
Pb-210	1.44E+12	Zr-95	8.96E+14
Pd-103	2.19E+15		

**Figure 1.** Representation of Voided Sphere Model

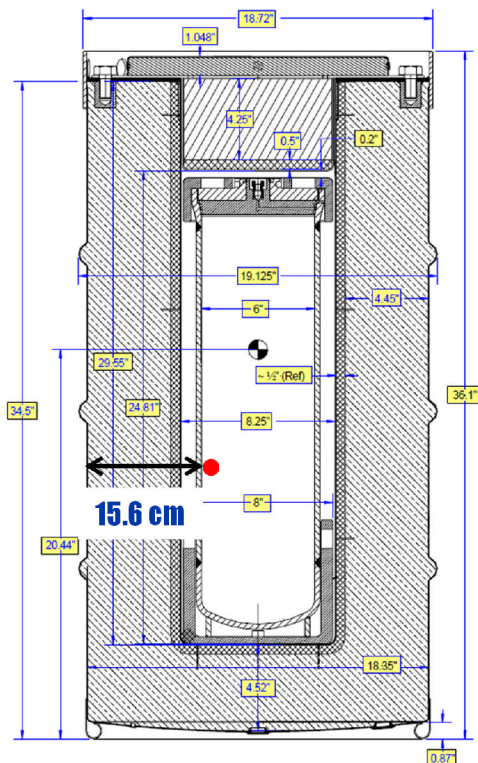


Figure 2. Diagram of Model 9977-96 Radioactive Material Transportation Package

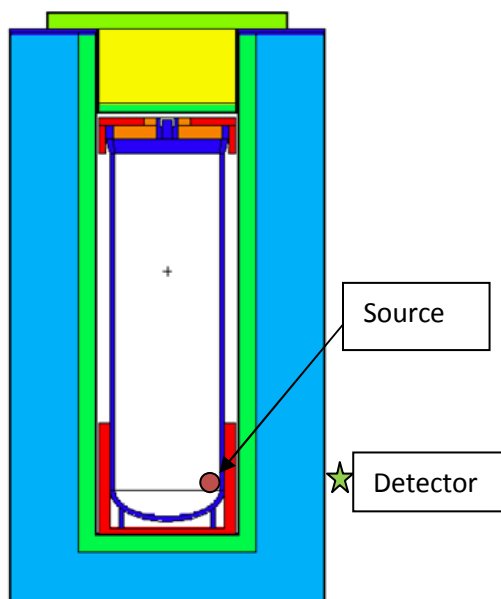


Figure 3. MCNP Input Model Used in the Shielding Calculations

The approach used to derive the results presented in this report, in terms of developing the source terms, the shielding model, and the calculations, are consistent with the guidance provided in Regulatory Guide 7.9, *Standard Format and Content of Part 71 Applications for Approval of*

Packages for Radioactive Material,^[9] and in the *Packaging Review Guide for Reviewing Safety Analysis Reports for Packaging*.^[10]

Results and Discussion

The isotopic masses which result in a dose rate of 190 mrem/hr at a distance of 15.6 cm using the voided sphere model are listed in Tables 4a and 4b. All sources are assumed to be point sources. The isotopic masses listed in Tables 4a and 4b were calculated assuming no shielding with only an air gap between the point source and the location of the assumed dose rate.

Table 4a. Limiting Isotopic Masses for Neutron Sources Using Voided Sphere Model

Isotopic Mass Limits for Voided Sphere Model [grams]: Distance=15.6 cm, Dose Rate=190 mrem/hr										
Isotope	Weight Percent Beryllium									
	0	0.1	0.5	1	5	10	30	50	70	90
Am-241	2.99E-03	2.99E-03	2.99E-03	2.99E-03	2.98E-03	2.98E-03	2.97E-03	2.97E-03	2.97E-03	2.96E-03
Am-243	2.78E-02	NA	NA	NA	NA	2.78E-02	2.78E-02	2.77E-02	2.77E-02	2.77E-02
Cf-252	1.92E-06	NA	NA	NA	NA	NA	NA	NA	NA	NA
Cm-244	8.96E-04	8.96E-04	8.95E-04	8.94E-04	8.90E-04	8.84E-04	8.68E-04	8.57E-04	8.50E-04	8.44E-04
Cm-248	1.12E-01	NA	NA	NA	NA	NA	NA	NA	NA	NA
Np-237	6.06E+00	NA	NA	NA	NA	6.05E+00	6.05E+00	6.05E+00	6.05E+00	6.05E+00
Pu-238	4.01E-03	4.00E-03	4.00E-03	4.00E-03	3.98E-03	3.96E-03	3.91E-03	3.87E-03	3.84E-03	3.82E-03
Pu-239	2.51E+00	2.51E+00	2.51E+00	2.51E+00	2.49E+00	2.47E+00	2.40E+00	2.36E+00	2.34E+00	2.32E+00
Pu-240	3.17E-01	3.17E-01	3.17E-01	3.17E-01	3.16E-01	3.15E-01	3.11E-01	3.08E-01	3.07E-01	3.05E-01
Pu-241	3.35E-03	NA	NA	NA	NA	3.35E-03	3.34E-03	3.34E-03	3.33E-03	3.33E-03
Pu-242	2.18E+01	2.18E+01	2.18E+01	2.18E+01	2.17E+01	2.16E+01	2.13E+01	2.12E+01	2.11E+01	2.10E+01

Table 4b. Limiting Isotopic Masses for Gamma Sources Using Voided Sphere Model

Isotopic Mass Limits for Voided Sphere Model [grams]: Distance=15.6 cm, Dose Rate=190 mrem/hr			
Isotope	Mass Limit [g]	Isotope	Mass Limit [g]
Ac-227	6.92E-05	Pm-147	5.74E-04
Cd-109	8.07E-06	Po-210	1.95E-01
Co-60	2.95E-06	Ra-226	2.98E-03
Cs-137	7.78E-05	Ru-106	2.86E-06
Eu-152	3.49E-05	Sc-46	1.18E-07
Fe-59	1.39E-07	Se-75	5.90E-07
Gd-153	7.56E-06	Sm-145	1.24E-05
Hf-181	6.51E-07	Sr-90	3.21E-04
Ho-166	4.70E-08	Tm-170	1.25E-05
Ir-192	8.05E-07	Yb-169	4.85E-07
Mn-54	1.19E-06	Zn-65	1.76E-06
Pb-210	2.80E-04	Zr-95	4.08E-07
Pd-103	1.74E-07		

The isotopic masses in the Model 9977-96 radioactive material transportation container that result in an external dose rate on the surface of the package of 190 mrem/hr are given in Tables 5a and 5b.

Table 5a. Limiting Isotopic Masses for Neutron Sources Using Model 9977-96 Radioactive Material Transportation Package

Isotopic Mass Limits for Model 9977 Packaging [grams]: Distance=15.6 cm, Dose Rate=190 mrem/hr										
Isotope	Weight Percent Beryllium									
	0	0.1	0.5	1	5	10	30	50	70	90
Am-241	1.65E+01	1.49E+01	1.08E+01	7.94E+00	2.86E+00	1.59E+00	6.96E-01	5.04E-01	4.20E-01	3.74E-01
Am-243	3.91E-01	NA	NA	NA	NA	3.87E-01	3.81E-01	3.77E-01	3.74E-01	3.72E-01
Cf-252	1.77E-06	NA	NA	NA	NA	NA	NA	NA	NA	NA
Cm-244	3.84E-01	3.53E-01	2.76E-01	2.29E-01	9.14E-02	5.28E-02	2.36E-02	1.72E-02	1.45E-02	1.28E-02
Cm-248	1.04E-01	NA	NA	NA	NA	NA	NA	NA	NA	NA
Np-237	7.97E+01	NA	NA	NA	NA	7.92E+01	7.86E+01	7.82E+01	7.79E+01	7.77E+01
Pu-238	4.18E+00	3.69E+00	2.51E+00	1.77E+00	5.97E-01	3.22E-01	1.41E-01	1.02E-01	8.46E-02	7.51E-02
Pu-239	1.16E+03	1.04E+03	7.62E+02	5.55E+02	2.02E+02	1.11E+02	4.91E+01	3.56E+01	2.97E+01	2.65E+01
Pu-240	2.67E+03	1.40E+03	4.84E+02	2.67E+02	6.39E+01	3.33E+01	1.39E+01	1.00E+01	8.14E+00	7.29E+00
Pu-241	2.03E+01	NA	NA	NA	NA	1.79E+00	7.83E-01	5.67E-01	4.73E-01	4.20E-01
Pu-242	2.22E+03	2.15E+03	2.10E+03	1.95E+03	1.51E+03	1.14E+03	6.70E+02	5.33E+02	4.64E+02	4.18E+02

Table 5b. Limiting Isotopic Masses for Gamma Sources Using Model 9977-96 Radioactive Material Transportation Package

Isotopic Mass Limits for Model 9977 Packaging [grams]: Distance=15.6 cm, Dose Rate=190 mrem/hr			
Isotope	Mass Limit [g]	Isotope	Mass Limit [g]
Ac-227	4.05E-04	Pm-147	3.31E-01
Cd-109	3.73E-02	Po-210	2.54E-01
Co-60	3.92E-06	Ra-226	6.15E-03
Cs-137	1.97E-04	Ru-106	9.93E-06
Eu-152	5.48E-05	Sc-46	1.57E-07
Fe-59	1.79E-07	Se-75	2.21E-06
Gd-153	2.32E-04	Sm-145	1.47E-02
Hf-181	1.25E-06	Sr-90	4.15E-03
Ho-166	4.30E-07	Tm-170	4.21E-04
Ir-192	1.39E-06	Yb-169	3.51E-06
Mn-54	1.56E-06	Zn-65	2.27E-06
Pb-210	1.93E-02	Zr-95	5.56E-07
Pd-103	1.33E-03		

The calculated limiting masses of the neutron-emitters using the voided sphere model (see Table 4a) ranged from 21 grams for Pu-242 to 1.92×10^{-6} grams for Cf-252. The limiting masses from largest to smallest were ordered as follows: Pu-242>Np-237>Pu-239>Pu-240>Cm-248>Am-243>Pu-238>Pu-241>Am-241>Cm-244>Cf-252.

The calculated limiting masses of gamma-emitters using the voided sphere model (see Table 4b) ranged from 0.195 grams for Po-210 to 4.70×10^{-8} grams for Ho-166. The limiting masses from largest to smallest were ordered as follows: Po-210>Ra-226>Pm-147>Sr-90>Pb-210>Cs-137>Ac-227>Eu-152>Tm-170>Sm-145>Cd-109>Gd-153>Co-60>Ru-106>Zn-65>Mn-54>Ir-192>Hf-181>Se-75>Yb-169>Zr-95>Pd-103>Fe-59>Sc-46>Ho-166.

As can be seen from examination of Table 4a, the allowable limiting masses of the neutron-emitters are rather insensitive to the beryllium concentration. This observation can be explained by the fact that the dose rates, and correspondently the limiting masses, of the neutron-emitters are governed mainly by their gamma emissions when using the voided sphere model. This is in contrast to the limiting masses for actinides with different beryllium concentrations when using the Model 9977-96 package (see Table 5a), where a significant amount of the gamma rays are attenuated by the packaging (i.e., stainless steel containment vessel, Celotex™, and stainless steel drum) and the dose rates (and limiting masses) are mainly governed by the neutron emissions.

The calculated limiting masses of the neutron-emitters using the Model 9977-96 package (see Table 5a) ranged from 2.67×10^3 grams for Pu-240 to 1.77×10^{-6} grams for Cf-252 for isotopes with no beryllium contamination. The limiting masses from largest to smallest were ordered as follows (no beryllium contamination): Pu-242>Np-237>Pu-239>Pu-240>Pu-241>Am-241>Am-243>Cm-248>Pu-238>Cm-244>Cf-252. Note that all of the neutron-emitting isotopes considered, except Cf-252, had limiting masses greater than 10mg when using the Model 9977-96 package. It should also be recognized that the limiting masses for Pu-239, Pu-240, and Pu-242 cannot be considered “small gram quantities” since they are all more than one kilogram.

Since the limiting masses for some isotopes were quite large, there was a concern that the assumption that the sources were *point sources* may not result in the most limiting mass. Additional analyses were performed whereby the sources were not assumed to be point sources but were assumed to be volume sources, according to their densities, and subcritical multiplication was taken into account. However, the self-shielding that occurs in the volume sources and the geometry effect of a volume source, more than compensated for the additional neutrons produced by subcritical multiplication, resulting in the point source models producing the most conservative values (i.e., lowest) for the limiting masses. It should be noted that these results are for homogeneous mixtures of beryllium and actinides and thus yield conservative mass limits. For actinides and beryllium combined in a different physical form, the mass limits could be higher.

The calculated limiting masses of gamma-emitters using the Model 9977-96 package (see Table 5b) ranged from 0.331 grams for Pm-147 to 1.57×10^{-7} grams for Sc-46. The limiting masses from largest to smallest were ordered as follows: Pm-147>Po-210>Cd-109>Pb-210>Sm-145>Ra-226>Sr-90>Pd-103>Tm-170>Ac-227>Gd-153>Cs-137>Eu-152>Ru-106>Co-60>Yb-169>Zn-65>Se-75>Mn-54>Ir-192>Hf-181>Zr-95>Ho-166>Fe-59>Sc-46.

It is of interest to determine the difference between masses calculated using the voided sphere model and those calculated when the package materials of the Model 9977-96 are taken into consideration. See Tables 6a and 6b for the fraction of allowable masses of the isotopes calculated with the Voided Sphere model to the Model 9977 package (i.e., in the Model 9977-97 package without and with the package materials present).

Table 6a. Fraction of Allowable Mass in Voided Sphere Model to Allowable Mass in the Model 9977-96 Radioactive Material Transportation Package for Neutron Emitters

Fraction of Allowable Mass in Voided Sphere Model to Allowable Mass in the Model 9977-96 Radioactive Material Transportation Package										
Isotope	Weight Percent Beryllium									
	0	0.1	0.5	1	5	10	30	50	70	90
Am-241	1.81E-04	2.00E-04	2.77E-04	3.76E-04	1.04E-03	1.88E-03	4.27E-03	5.89E-03	7.06E-03	7.92E-03
Am-243	7.11E-02	NA	NA	NA	NA	7.19E-02	7.29E-02	7.37E-02	7.42E-02	7.46E-02
Cf-252	1.08E+00	NA	NA	NA	NA	NA	NA	NA	NA	NA
Cm-244	2.33E-03	2.54E-03	3.24E-03	3.90E-03	9.74E-03	1.67E-02	3.68E-02	4.98E-02	5.86E-02	6.59E-02
Cm-248	1.07E+00	NA	NA	NA	NA	NA	NA	NA	NA	NA
Np-237	7.60E-02	NA	NA	NA	NA	7.64E-02	7.70E-02	7.73E-02	7.76E-02	7.78E-02
Pu-238	9.58E-04	1.08E-03	1.60E-03	2.26E-03	6.68E-03	1.23E-02	2.78E-02	3.81E-02	4.55E-02	5.09E-02
Pu-239	2.17E-03	2.41E-03	3.30E-03	4.52E-03	1.23E-02	2.23E-02	4.90E-02	6.64E-02	7.86E-02	8.75E-02
Pu-240	1.19E-04	2.27E-04	6.55E-04	1.19E-03	4.94E-03	9.44E-03	2.24E-02	3.09E-02	3.77E-02	4.19E-02
Pu-241	1.65E-04	NA	NA	NA	NA	1.87E-03	4.27E-03	5.88E-03	7.05E-03	7.92E-03
Pu-242	9.81E-03	1.01E-02	1.04E-02	1.11E-02	1.44E-02	1.89E-02	3.19E-02	3.97E-02	4.54E-02	5.01E-02

Table 6b. Fraction of Allowable Mass in Voided Sphere Model to Allowable Mass in the Model 9977-96 Radioactive Material Transportation Package for Gamma Emitters

Fraction of Allowable Mass in Voided Sphere Model to Allowable Mass in the Model 9977-96 Radioactive Material Transportation			
Isotope	Fraction	Isotope	Fraction
Ac-227	1.71E-01	Pm-147	1.74E-03
Cd-109	2.17E-04	Po-210	7.70E-01
Co-60	7.54E-01	Ra-226	4.84E-01
Cs-137	3.94E-01	Ru-106	2.88E-01
Eu-152	6.37E-01	Sc-46	7.48E-01
Fe-59	7.80E-01	Se-75	2.67E-01
Gd-153	3.26E-02	Sm-145	8.40E-04
Hf-181	5.23E-01	Sr-90	7.74E-02
Ho-166	1.09E-01	Tm-170	2.98E-02
Ir-192	5.81E-01	Yb-169	1.38E-01
Mn-54	7.65E-01	Zn-65	7.74E-01
Pb-210	1.45E-02	Zr-95	7.33E-01
Pd-103	1.31E-04		

An examination of the fraction of allowable mass from the voided sphere model to allowable mass in the Model 9977-96 radioactive material transportation package for neutron emitters [excluding Cm-248 & Cf-252] (Table 6a) shows that the fractions ranged from 1.19×10^{-4} to 8.75×10^{-2} . These data show that the results from the voided sphere model, for most neutron emitting isotopes, are conservative relative to the limiting masses for the Model 9977-96 package. For the notable exceptions, which are Cm-248 and Cf-252, the results from the voided sphere model are non conservative, by 7% and 8% respectively, relative to the limiting masses for these isotopes in the Model 9977-96. These seemingly non-intuitive results can be explained as follows. Except for these two isotopes, all the other neutron emitters have gamma to neutron source ratios that range from hundreds to tens of thousands. The vast majority of the gammas are in the low energy range that are attenuated by the materials in the Model 9977-96 package

whereas the neutrons are not attenuated to the same extent. Thus while the neutron contribution to the dose rate is always higher for the source inside the Model 9977-96 package when compared to the void case, the gamma contribution to the dose rate is dominant for the void case compared to that when the source is inside the package. As a result, the mass limit for the void case is lower than for the Model 9977-96 case. For Cm-248 and Cf-252, which are both large spontaneous fission neutron sources, the gamma to neutron source ratios are 5.5 and 4.2, respectively. Thus, in these cases the neutron dose rate is the dominant contributor to the total dose rate in both the void and packaged cases. In addition, the neutron scattering enhances the dose rate at the package surface leading to a lower mass limit for the Model 9977-96 case compared to the void case.

Since the mass limit results for Cm-248 and Cf-252 obtained using the voided sphere model are non-conservative relative to the mass limit results for these isotopes in the Model 9977-96 package, it is suggested that for application of the voided sphere mass limits for Cm-248 and Cf-252 to other transportation packages that the mass limits be reduced by about 10%. This would ensure that the 5% conservative margin for the results obtained by assuming a dose rate of 190 mrem/hr is retained (recall that the regulatory dose rate limit on the external surface of the transportation package is 200mrem/hr).

When the fractions of allowable masses from the voided sphere model to allowable masses in the Model 9977-96 radioactive material transportation package for gamma emitters are examined (Table 6b), it is seen that the fractions are generally much high than those of the neutron emitters and range from 1.31×10^{-4} to 0.78, with 68% of the gamma emitting isotopes considered having a fraction greater than 0.1. For the gamma emitting isotopes, the relative conservatism of the results from the voided sphere model to the Model 9977-96 package is generally due to the primary gamma energy or energies emitted by the isotope. Lower energy gamma rays are appreciably attenuated by the packaging materials in the Model 9977-96 and therefore the results from the voided sphere model for those cases are more conservative. In general, the gamma-emitting isotopes that primarily emit higher energy gamma rays will be identified by higher fractions in Table 6b. The differences in the mass ratios for some of the harder spectrum gamma emitters is not statistically significant. For example, the results show the Co-60 with a harder spectrum has a ratio smaller than that of Fe-59, where it would be expected to have a slightly higher value.

It should be noted that the limiting isotopic masses presented in this document for the Model 9977-96 package are based solely on external radiation level considerations, and that the actual limiting mass for transportation in this packaging under the restrictions of 10 CFR 71 and the Model 9977-96 Safety Analysis Report for Packaging (SARP) may be based on other considerations, such as criticality, mass, volume, and heat generation etc.. The Model 9977-96 mass limits for both gamma emitters and neutron emitters are also applicable to Models 9978 and 9975 packagings noting that these mass limits will be on the conservative side for these two packagings.

Applicability

The isotopic limiting mass results from the voided sphere model are widely applicable for general shielding considerations and to radioactive material packaging in particular. The voided sphere results can be conservatively applied (except the results for Cm-248 and Cf-252) to essentially any transportation and storage package since the model assumes no shielding is present.

Additionally, the voided sphere results can be applied to shielding considerations as related to facility operations. The voided sphere data presented in this paper can be used as a reference for dose rates for unshielded sources, where the assumption of a point source is valid.

For application of the voided sphere results to a distance other than 15.6 cm or to a dose rate criterion of other than 190 mrem/hr, the limiting mass would need to be scaled:

$$M_{LP} = \left(\frac{D_C}{190} \right) \left(\frac{d_p}{15.6} \right)^2 M_{L,15.6} \quad (E1)$$

where:

- M_{LP} is the isotopic mass limit [grams],
- d_p is the distance for the package under consideration (usually the minimum distance from the inner wall of the containment (or confinement vessel) [cm],
- $M_{L(15.6)}$ is the isotopic mass limit calculated using the Voided Sphere model at a distance of 15.6 cm [grams] that produces a dose rate of 190 mrem/hr, and
- D_C is the dose rate criterion of interest [mrem/hr].

For mixtures of isotopes, the sum of the fraction of each isotope to its individual limit should be less than or equal to 1, i.e.

$$\sum_{i=1}^n \frac{M_i}{M_{LP,i}} \leq 1 \quad (E2)$$

where:

M_i is the mass of the i^{th} isotope in the mixture containing n isotopes and $M_{LP,i}$ is the maximum allowed mass for that isotope that may be calculated using equation E1. Equation E2 is applicable to both the voided and the Model 9977-96 cases, where M_{LP} is obtained from Tables 5a and 5b (equation E1 is not applicable to the Model 9977-96).

Conclusions and Future Work

This report presents a set of mass limits for commonly used sources and other material found at various DOE sites that can be shipped in unshielded containers or the 9977, 9978, and 9975

packagings. For the case of common actinides that contain mixed in light elements, a conservative set of limits has been presented based on various levels of beryllium impurity.

This work could be extended in many aspects, including: (1) consideration of additional isotopes, such as additional medical and industrial isotopes, (2) additional transportation package-specific calculations, such as calculations of limiting isotopic masses for materials in the shielded inserts for the Model 9977-96, and (3) generic shielding calculations, where a given thickness of shielding, such as lead or polyethylene, is considered when determining the allowable mass limits. Examples of other medical, industrial, or rarely-shipped isotopes that could be considered in future work.

The Model 9977-96 package can be used with three different shielded containers or inserts: (1) the SGQ-SC1, which is made of lead and encapsulated in stainless steel with a threaded stainless steel closure, (2) the SGQ-SC2, which is constructed of high-density polyethylene (HDPE), and (3) the SGQ-SC3, which is constructed of tungsten and is also encapsulated in stainless steel. Additional calculations of isotopic mass limits in the Model 9977-96 could include considerations of the lead and tungsten shielded inserts for isotopes that are gamma emitters, and consideration of the HDPE shielded insert for isotopes (and actinides with light element impurities) that are primarily neutron emitters.

Moreover, other future work could include developing a generalized approach for light element impurities other than beryllium. Although, use of the beryllium impurity concentrations for determining isotopic mass limits for other light element impurities is conservative, the mass limit envelope for alpha-emitting isotopes with these other light element impurities could be broadened and therefore optimize shipments. Since the alpha-neutron (i.e., α -n) behavior of light elements relative to beryllium is known, a generalized approach for determining the α -n behavior of alpha-emitting isotopes with light element impurities, including mixtures of isotopes with mixtures of light element impurities, could be developed.

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